Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- (Currently amended)
 An anticancer or an anti-metastatic agent for gene therapy containing A pharmaceutical composition for treating a solid tumor, or metastasis thereof, said composition comprising a gene carrier or cells harboring a gene encoding a recombinant protein consisting of human apolipoprotein(a) kringle KIV9-KIV10-KV (LK68) or KV (LK8) gene as an effective ingredient.
- (Currently amended) The agent composition according to claim 1, wherein the LK68 gene comprises a nucleotide sequence represented by SEO, ID, No. 1.
- 3. (Currently amended) The agent composition according to claim 1, wherein the gene carrier harboring the LK68 gene is a vector or a recombinant virus.
- 4. (Currently amended) The agent composition according to claim 3, wherein the vector is selected from a group consisting of a linear DNA vector, a plasmid DNA vector and a recombinant viral vector.
- 5. (Currently amended) The agent composition according to claim 3, wherein the recombinant virus is selected from a group consisting of retrovirus, adenovirus, adenovassociated virus, herpes simplex virus and lentivirus.
- 6. (Currently amended) The agent composition according to claim 1, wherein the cells are selected from a group consisting of hematopoictic stem cells, dendritic cells, autologous tumor cells and established tumor cells.

- 7. (Currently amended) The agent composition according to claim 1, wherein the gene carrier is selected from a group consisting of pSecTag-LK68, pLXSN-LK68, rAAV-LK68 and pAAV-LK68.
- 8. (Currently amended) The agent composition according to claim 1, wherein the LK8 gene comprises a nucleotide sequence represented by SEQ. ID. No. 2.
- 9. (Currently amended) The agent composition according to claim 1, wherein the gene carrier harboring the LK8 gene is a vector or a recombinant virus.
- 10. (Currently amended) The agent composition according to claim 9, wherein the vector is selected from a group consisting of a linear DNA vector, a plasmid DNA vector and a recombinant viral vector.
- 11. (Currently amended) The agent composition according to claim 9, wherein the recombinant virus is selected from a group consisting of retrovirus, adenovirus, adenovirus associated virus, herpes simplex virus and lentivirus.
- 12. (Currently amended) The agent composition according to claim 9, wherein the gene carrier is selected from a group consisting of pSecTag-LK8, pLXSN-LK8, rAAV-LK8 and pAAV-LK8.
- 13. (Currently amended) The $\frac{\text{agent composition}}{\text{according to claim 3, wherein the vector is included by 0.05 <math>\sim$ 500 mg.
- 14. (Currently amended) The $\frac{\text{agent composition}}{10^{12}}$ according to claim 3, wherein the recombinant virus is included by 10^3 10^{12} IU.
- 15. (Currently amended) The $\frac{\text{agent composition}}{\text{according to claim 1, wherein the}}$ tells are included by 10^3 10^8 e.a.

- 16. (Currently amended) The agent composition according to claim 1, wherein the eaneer solid tumor is selected from a group consisting of colon carcinoma, liver cancer, lung cancer, breast cancer, brain tumor, prostatic carcinoma, skin cancer, stomach cancer, pancreas cancer, lymphoma, kidney cancer, ovarian cancer and metastatic tumor.
- 17. (Currently amended) The agent composition according to claim 16, wherein the cancer solid tumor is selected from a group consisting of colon carcinoma, liver cancer, lymphoma of and metastatic tumor.
- 18. (Currently amended) A method for the prevention or the treatment of a solid tumor, which includes a step of parenteral administration of the agent for gene therapy composition of claim 1 to an individual.
- 19. (Currently amended) The method according to claim 18, wherein the prevention or the treatment of a solid tumor is accomplished by the inhibition of the growth and the metastasis of the solid tumor.
- 20. (Currently amended) The method according to claim 18, wherein the administration of a gene carrier harboring human apolipoprotein(a) kringle KIV9-KIV10-KV (LK68) or KV (LK68) gene is accomplished by a method selected from a group consisting of chemical method, physical method, conjugation using liposome, a method using receptor and virus-etc.
- 21. (Currently amended) The method according to claim 18, wherein the administration is characterized by injecting cells selected from a group consisting of hematopoietic stem cells, dendritic cells, autologous tumor cells and established tumor cells transfected with human apolipoprotein(a) kringle KIV9-KIV10-KV(LK68) or KV(LK8) gene to a patient.

- 22. (Currently amended) The $\frac{\text{agent composition}}{\text{composition}}$ according to claim 9, wherein the vector is included by $0.05 \sim 500$ mg.
- 23. (Currently amended) The $\frac{\text{agent composition}}{10^{12}}$ according to claim 9, wherein the recombinant virus is included by 10^3 10^{12} IU.